### Appendix E-2

"HAP Benefits Analysis in Section 812 Reports to Congress; Briefing for SAB/EPA Workshop, June 22, 2000,"

Presented by Mr. James DeMocker

### HAP Benefits Analysis in §812 Reports to Congress

Briefing for SAB/EPA Workshop June 22, 2000



## §812 Benefit-Cost Analyses Analytical Requirements

- "(a)...The Administrator shall conduct a comprehensive analysis of this Act on the public health, economy, and environment... [which] should consider the costs, benefits and other effects...[of] each standard issued for... (2) a hazardous air pollutant listed under §112, including any technology-based standard and any risk-based standard..."
- "(b)...The Administrator shall assess how benefits are measured in order to assure that damage to human health and the environment is more accurately measured and taken into account."

## §812 Benefit-Cost Analyses Review Requirements

• "(f)...The Administrator shall appoint an Advisory Council... [consisting of] recognized experts in... health and environmental effects of air pollution, economic analysis, environmental sciences, and other [appropriate] fields."

• "(g)...The Council shall review... the data... the methodology... and the findings of such report, and make recommendations to the Administrator concerning the validity and utility of such findings."

### §812 Benefit & Cost Estimation

- "Retrospective Study"
  - Submitted to Congress October 1997
  - Direct costs aggregated and fed to macro model
  - Benefits by pollutant as data and models allowed
- "Prospective Study"
  - Submitted to Congress November 1999
  - Direct costs estimated by title / major provision
  - Benefits by pollutant as data and models allowed

# Retrospective Study Stationary Source Pollutants 14 key HAPs from Cancer Risk Study (1990):

- arsenic
- asbestos
- benzene
- 1,3-butadiene
- carbon tetrachloride
- chloroform
- chromium (VI)

- dioxin
- ethylene dichloride
- ethylene dibromide
- formaldehyde
- gasoline vapors
- product of incomplete combustion (PICs)
- vinyl chloride

### Retrospective Study Stationary Source Method

Incidence change assumed proportional to emissions change

$$I_{ty} = I_{by} \times \frac{A_{ty}}{A_{by}} \times \frac{P_{ty}}{P_{by}} \times \frac{(1-C_{ty})}{(1-C_{by})}$$

```
I = incidence (from CRS) by = base year (85)

A = activity (from macro model) ty = target year (70, 75, 80, 90)

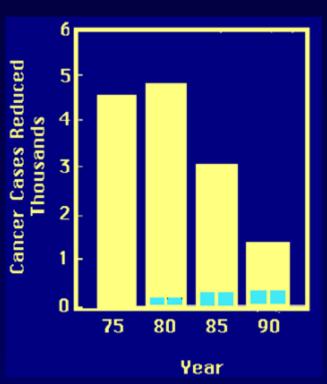
D = normalistics
```

P = population

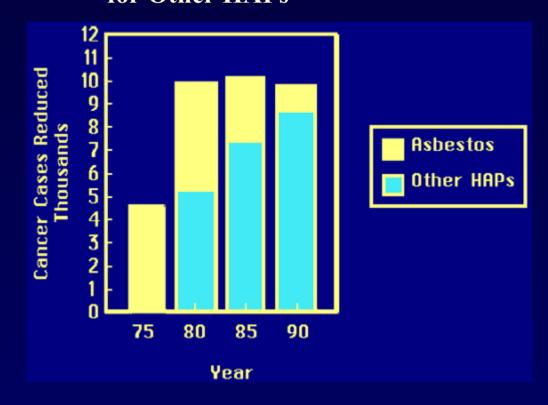
 $C = control\ efficiency\ (from\ CTGs,\ BIDs,\ regs,\ experts)$ 

## Retrospective Study Stationary Source Findings

**Lower Bound for Other HAPs** 



**Upper Bound for Other HAPs** 



## Retrospective Study Stationary Source Review Issues

- Estimated incidence for vinyl chloride and asbestos much higher than historical incidence
- Cancer Risk Study designed for only rough order-ofmagnitude estimates
  - Unit risk factors are upper-bound estimates
  - Exposure estimates are typically upper-bound (MEI)
- Control efficiencies assumed uniform across facilities and 100% compliance with regulations

## Retrospective Study Report to Congress Presentation

- HAP benefits excluded from primary analysis described in Appendix
- Quantitative analyses with caveats
  - Stationary source cancer incidence reduction estimates
  - Motor vehicle exposure reduction estimates
- Qualitative discussions
  - non-cancer health effects
  - ecosystem effects

#### -- Health Research Recommendations --

- Address additional pollutants
- Address mechanisms with pharmacokinetics
- Address variations in human susceptibility
- Address interactive effects of multiple exposures
- Develop alternatives to cancer upper-bound methods
- Develop D/R relationships for non-cancer effects
- Develop methods for acute exposure effects

- -- Exposure Research Recommendations --
- Expand data collection: control efficiencies, HAP speciation, facility locations and operating parameters
- Develop more comprehensive exposure models
- Refine uncertainty analysis methods

#### -- Ecosystem Research Recommendations --

- Estimate levels of bioaccumulating toxics in media
- Correlate levels of bioaccumulating toxics with exposures, concentrations, and adverse effects
- Develop wildlife correlate to RfD or D/R relationship
- Address effects of mixtures
- Address additional ecosystems
- Address wetland species and functions

-- Valuation Research Recommendations --

 Address additional endpoints consistent with kinds of damages expected

• Initiate broad-scope economic valuation using survey techniques

# Prospective Study Methodology Alternatives Presented -- National Scale --

- Assessment System for Population Exposure Nationwide (ASPEN)
  - Emissions inventory
    - multiple pollutants
  - Air dispersion model
    - point, area, and mobile source categories
  - Exposure model (not completed)

# Prospective Study Methodology Alternatives Presented -- National Scale --

- Advantages
  - Includes treatment of
    - reactive decay (simplified)
    - secondary formation (simplified)
    - long-range transport (continental scale)
    - wet and dry deposition (parameterized)
  - Emissions/Dispersion well documented:
    - sensitivity analysis
    - model performance evaluation
    - uncertainty analyses

# Prospective Study Methodology Alternatives Presented -- National Scale --

#### Limitations

- National emission inventory uncertainties
- Gaussian model limitations
- Meso-scale transport not addressed (50 200 km)
- Re-suspension not addressed
- Not stochastic
- Spatial and temporal peaks not addressed
- Indoor sources not addressed
- Indirect exposures not addressed

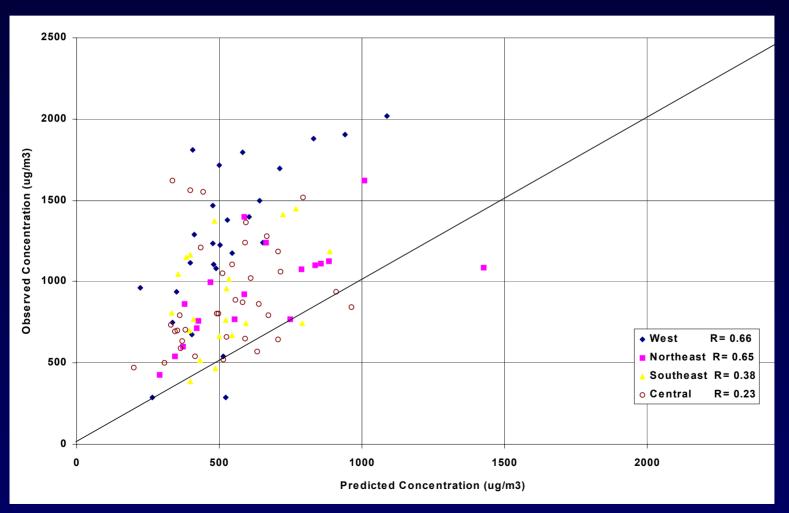
# Prospective Study Methodology Alternatives Presented -- Local Scale / Case Study --

- Air Quality Integrated Management System (AIMS)
- Developed for Baltimore and planned for Houston and Chicago
- Integrates routinely collected data (measured air quality, emissions, and meteorological data) and dispersion modeling

### Prospective Study Review Issues Raised

- Resources for in-depth analysis for 188 HAPs prohibitive: find priority HAPs
- Unit Risk Factors are upper-bound estimates
- Limited ambient monitoring data to validate ambient concentration estimates
- Exposure assessment limitations
  - 50 km downwind distance for dispersion
  - lack of attention to indirect pathways (e.g., Hg, dioxin)
  - ASPEN preliminary performance evaluation concerns

## ASPEN Model Performance 1990 Carbon Monoxide



## Prospective Study: Report to Congress Presentation

- No quantified benefits
- Expect benefits from MACT and incidental to criteria pollutant control
- Besides cancer inhalation impacts, other potential benefits include reductions in:
  - Non-cancer health effects
  - Indirect non-inhalation exposure
  - Ecological and welfare effects

## Prospective Study Report to Congress Research Recommendations

- Workshops to address HAP benefits challenges:
  - toxicology/risk assessment
  - exposure assessment
  - economics
- Investigate use of EPA's Air Toxics Data Archive of measurement data from state / local programs
- Explore whether "supersite" monitoring programs can provide HAP ambient concentration data

### Future §812 Studies

• Pondering potential scope, objectives, and reference period for "812 III"

• Detailed analytic blueprint to be developed, and HAP Workshop outcomes will be considered

• SAB Council and HEES will be asked to review analytical blueprint prior to initiation of work

### Backup slides

### Retrospective Study: Mobile Source Analysis

-- Methods --

- Based on Motor Vehicle Related Air Toxics Study (1993)
- Exposure estimated for CO using measured concentrations and HAPEM-MS
- Exposure to HAPs assumed proportional to emission factors

### Retrospective Study: Mobile Source Analysis

#### -- Methods and Data--

$$E = ((A \times C) - B) \times M \times S \times \frac{VOC \times HAP}{CO}$$

E = exposure concentration

A = annual average CO ambient concentration (AIRS)

C = CO ambient to CO exposure concentration ratio (HAPEM)

B = CO background concentration (reported measurements)

M = fraction of CO emissions from mobile sources

S = scenario-to-control scenario CO emission factor ratio

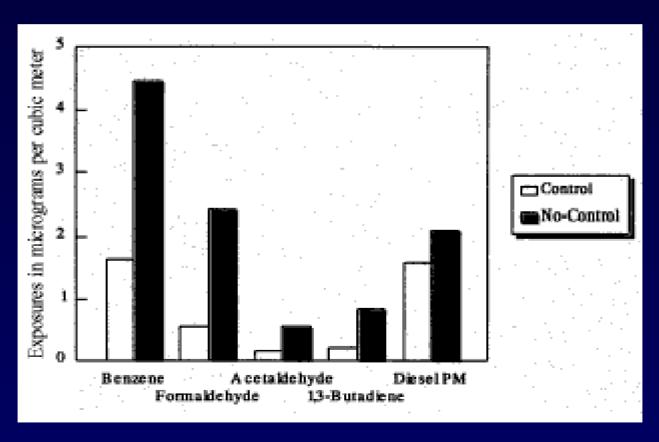
*VOC* = VOC mobile emission factor by, scenario/year

HAP = HAP speciation factor for mobile source VOC, by scenario/year

*CO* = mobile source emission factor, by scenario/year

## Retrospective Study: Mobile Source Analysis

-- Findings --



## Future 812 Studies Tools Needed

- Expanded air toxics monitoring data
  - 90 new monitors by end of FY00
  - Air Toxics Data Archive to supplement AIRS with state and local data
- Improved emissions inventories
  - 1996 National Toxics Inventory (NTI)
- Evaluation/enhancement of air quality and exposure modeling tools
- Expanded risk data and improved methods

- Current risk assessment state-of-the-art
  - Probabilistic estimates for cancer
  - Reference doses/concentrations for non-cancer
    - More sophisticated D/R assessments for some criteria pollutants
  - Mixtures
    - Sum of upper-bounds for cancer
    - Hazard index for non-cancer

• Recent trends in risk assessment

- Cancer: mix of probabilistic (no threshold) and reference concentrations (threshold)
- Non-cancer: modeling and distributional approaches
- Dosimetry models focused on tissue concentrations

#### Potential sources of bias in risk estimates

- Linear high-to-low dose extrapolation
- Cross species scaling factor
- Treatment of untested chemicals and other data gaps
- Latent effects
- Use of most sensitive test results
- Non-cancer uncertainty factors
- Magnitude and severity of effects
- Route-to-route extrapolation
- Benchmark response rate (LED10 instead of NOAEL)
- Additive treatment of mixtures

- Uncertainty in risk estimates
  - Types
    - Causal link between exposure and effects
    - Magnitude of risk
  - Can use analysis of quantifiable uncertainty to develop central risk estimate
  - Unquantifiable uncertainty may still lead to bias
    - use of sensitive species
    - consideration of non-relevant effects

- Topics for discussion
  - How to characterize a distribution of risk estimates as an input to benefits assessment
  - How to characterize the value of reducing exposure in a reference dose framework: proportion of people above RfD, contingent valuation, other?
  - How to characterize benefits when uncertainty is great: point estimate, range, other?
  - Are some benefits better left unquantified?